PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

(Chapter II of the Patent Cooperation Treaty)

(PCT Article 36 and Rule 70)

REC'D 3 0 MAR 2006

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Applicant's or agent's file reference PC32214A	FOR FURTHER ACTION	See Form PCT/IPEA/416						
International application No. PCT/IB2005/001140	International filing date (daytmon 25.04.2005	th/year) Priority date (day/month/year) 06.05.2004						
International Patent Classification (IPC) or national classification and IPC INV. C07D207/16 C07D211/26 C07D211/60 C07D211/96 C07D265/30 A61K31/401 A61K31/445 A61K31/5375								
Applicant PFIZER INC. et al.								
This report is the international prel Authority under Article 35 and tran		tablished by this International Preliminary Examining ling to Article 36.						
2. This REPORT consists of a total of	of 8 sheets, including this cover	r sheet.						
3. This report is also accompanied by	y ANNEXES, comprising:							
a. sent to the applicant and to	the International Bureau) a tot	al of sheets, as follows:						
sheets of the description, claims and/or drawings which have been amended and are the basis of this report and/or sheets containing rectifications authorized by this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions).								
	beyond the disclosure in the international application as filed, as indicated in item 4 of Box No. I and the							
sequence listing and/or tab		ype and number of electronic carrier(s)) , containing a ic form only, as indicated in the Supplemental Box inistrative instructions).						
4. This report contains indications re	lating to the following items:							
☐ Box No. I Basis of the repo	ort							
☐ Box No. II Priority	-							
⊠ Box No. III Non-establishme	ent of opinion with regard to no	velty, inventive step and industrial applicability						
☐ Box No. IV Lack of unity of i	invention							
	Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement							
☑ Box No. VI Certain docume	nts cited							
☐ Box No. VII Certain defects i	☐ Box No. VII Certain defects in the international application							
☐ Box No. VIII Certain observations on the international application								
Date of submission of the demand	Date of	completion of this report						
20.06.2005	28.03	.2006						
Name and mailing address of the International preliminary examining authority:	al Authori	zed officer						
European Patent Office D-80298 Munich Rudolf, M								
Tel. +49 89 2399 - 0 Tx: 52365 Fax: +49 89 2399 - 4465	56 epmu d							
	Teleph	one No. +49 89 2399-						

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

International application No. PCT/IB2005/001140

	Box No. I Ba	asis of the report			
1.		vith regard to the language, this report is based on the international application in the language in which it was led, unless otherwise indicated under this item.			
	which is th ☐ interna ☐ publica	t is based on translations from the original language into the following language, ne language of a translation furnished for the purposes of: tional search (under Rules 12.3 and 23.1(b)) ation of the international application (under Rule 12.4) tional preliminary examination (under Rules 55.2 and/or 55.3)			
2.	With regard to the elements* of the international application, this report is based on (replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report):				
	Description, Pa	iges .			
	1-147	as originally filed			
	Claims, Numbe	are			
	1-15	as originally filed			
		and on growing models.			
	☐ a sequence	ce listing and/or any related table(s) - see Supplemental Box Relating to Sequence Listing			
3.	☐ The amen	dments have resulted in the cancellation of:			
	☐ the des☐ the cla	scription, pages			
	☐ the dra	wings, sheets/figs			
	☐ the sec	quence listing <i>(specify)</i> : ple(s) related to sequence listing <i>(specify)</i> :			
4.	had not been r Supplemental	t has been established as if (some of) the amendments annexed to this report and listed below made, since they have been considered to go beyond the disclosure as filed, as indicated in the Box (Rule 70.2(c)).			
	⊔ the des □ the cla	scription, pages ims. Nos.			
	☐ the dra	awings, sheets/figs			
		quence listing <i>(specify)</i> : ple(s) related to sequence listing <i>(specify)</i> :			
	•	4 applies, some or all of these sheets may be marked "superseded."			

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

International application No. PCT/IB2005/001140

Box No. III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability				
The	the questions whether the claimed invention appears to be novel, to involve an inventive step (to be non- povious), or to be industrially applicable have not been examined in respect of:			
	the entire international application,			
	claims Nos. 14,15 with respect to industrial applicability			
	because:			
⊠	the said international application, or the said claims Nos. 14, 15 with respect to industrial applicability relate to the following subject matter which does not require an international preliminary examination (specify):			
	see separate sheet	•		
	the description, claims or drawings (indicate particular elements below) or said claims Nos. are so unclear that no meaningful opinion could be formed (specify):			
	the claims, or said claims Nos. are so inadequately supported by the description that no meaningful opinion could be formed.			
	no international search report has been established for the said claims Nos.			
	the nucleotide and/or amino acid sequence listing does not comply with the standard provided for in Annex C of the Administrative Instructions in that:			
	the written form		has not been furnished	
			does not comply with the standard	
,	the computer readable form		has not been furnished	
			does not comply with the standard	
	the tables related to the nucleo not comply with the technical re	tide a equire	and/or amino acid sequence listing, if in computer readable form only, do ements provided for in Annex C-bis of the Administrative Instructions.	
	See separate sheet for further	detai	is	

Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)

Yes: Claims

11,12

No: Claims 1-10,13-15

Inventive step (IS)

Claims

11,12

Claims No:

1-10, 13-15

Industrial applicability (IA)

Yes: Claims

1-13

Claims No:

2. Citations and explanations (Rule 70.7):

see separate sheet

Certain documents cited Box No. VI

1. Certain published documents (Rule 70.10) and /or

2. Non-written disclosures (Rule 70.9)

see separate sheet

Re Item III.

Claims 14-15 relate to subject-matter considered by this Authority to be covered by the provisions of Rule 67.1(iv) PCT. Consequently, no opinion will be formulated with respect to the industrial applicability of the subject-matter of these claims (Article 34(4)(a)(I) PCT).

Re Item V.

- 1. Reference is made to the following documents:
 - D1: WO 00/33788 A (AMERICAN BIOGENETIC SCIENCES INC; UNIVERSITY COLLEGE DUBLIN; SZMUSZKOV) 15 June 2000 (2000-06-15)
 - D2: BIN HO, PRABHA M. VENKATARANGAN, SHARON F. CRUSE ET AL.: "Synthesis of 2-piperidinecarboxylic acid derivatives as potential anticonvulsants" EUR. J. MED. CHEM., vol. 33, 1998, pages 23-31, XP002348704
 - D3: MIN SHEN, ARNAUD LETIRAN, YUNDE XIAO, ALEXANDER GOLBRAIKH, HAROLD KOHN, ALEXANDER TROPSHA: "Quantitative Structure-Activity Relationship Analysis of Functionalized Amino Acid Anticonvulsant Agents Using k Nearest Neighbor and Simulated Annealing PLS Methods" J. MED. CHEM., vol. 45, 2002, pages 2811-2823, XP002348705
 - D4: US 3 931 139 A (WISSMANN ET AL) 6 January 1976 (1976-01-06)
 - D5: BOULOS ZACHARIE, NANCIE MOREAU, CHRISTOPHER DOCKENDORFF: "A Mild procedure for the Reduction of Pyridine N-Oxides to Piperidines Using Ammonium Formate" J. ORG. CHEM., vol. 66, 2001, pages 5264-5265, XP002348706
 - D6: WO 02/18335 A (YAMANOUCHI PHARMACEUTICAL CO., LTD; TORAY INDUSTRIES, INC; MORIHIRA, K) 7 March 2002 (2002-03-07)
 - D7: VIDYADHAR S. RANADE, ROEL PRINS: "Diastereoselective Hydrogenation of (S)-Proline-2-methylanilide" JOURNAL OF CATALYSIS, vol. 185, 1999, pages 479-486, XP002348707
 - D8: TSUTOMU MIMOTO, RYOHEI KATO, HARUO TAKAKU, SATOSHI NOJIMA, KEISUKE TERASHIMA ET AL.: "Structure-Activity Relationship of Small-Sized

- HIV Protease Inhibitors Containing Allophenylnorstatine" J.MED. CHEM., vol. 42, 1999, pages 1789-1802, XP002348708
- D9: STEVEN K. DAVIDSEN, PAUL D. MAY, JAMES B. SUMMERS: "Di-tert-butyl N-Acylimidodicarbonates as Isolable Acylating Agents: Mild Conversion of Primary Carboxamides to Substituted Amides" J. ORG. CHEM., vol. 56, 1991, pages 5482-5485, XP002348709
- D10: JOSEPH L. DUFFY, NANCY J. KEVIN, BRIAN A. KIRK, KEVIN T. CHAPMAN, WILLIAM A. SCHLEIF ET AL.: "Synthesis and Activity of Novel HIV Protease Inhibitors with Improved Potency Against Multiple PI-Resistant Viral Strains" BIOORGANIC &; MEDICINAL CHEMISTRY LETTERS, vol. 12, 2002, pages 2423-2426, XP002348710
- D11: JU YOUNG LEE, YONG JUN CHUNG, YOE-SIK BAE, SUNG HO RYU, BYEANG HYEAN KIM: "Synthesis of hexapeptide and tetrapeptide analogues of the immunomodulating peptides" J. CHEM. SOC. PERKIN TRANS. 1, 1998, pages 359-365, XP002348711
- D12: KOICHI KAWASAKI, KATSUHIKO HIRASE, MASANORI MIYANO, TOSHIKI TSUJI, MASANORI IWAMOTO: "Amino Acids and Peptides XVI. Synthesis of N-Terminal Tetrapeptide Analogs of Fibrin a-Chain and Their Inhibitory Effects on Fibrinogen/Thrombin Clotting" CHEM. PHARM. BULL., vol. 40, no. 12, 1992, pages 3253-3260, XP001207622
- D13: HAE YOON RHYOO, YOUNG-AE YOON, HEE-JUNG PARK, YOUNG KEUN CHUNG: "Use of amino amides derived from proline as chiral ligands in the ruthenium(II)-catalyzed transfer hydrogenation reaction of ketones" TETRAHEDRON LETTERS, vol. 42, 2001, pages 5045-5048, XP002348712
- D14: SHU KOBAYASHI, HIROMI UCHIRO, YUKO FUJISHITA, ISAMU SHIINA, TERUAKI MUKAIYAMA: "Asymmetric Aldol Reaction between Achiral Silyl Enol Ethers and Achiral Aldehydes by Use of a Chiral Promoter System" J. AM. CHEM. SOC., vol. 113, 1991, pages 4247-4252, XP002348776
- D15: MAKOTO NAKAJIMA, IRIE MIYOSHI, KUMIKO KANAYAMA, SHUN-ICHI HASHIMOTO: "Enantioselective Synthesis of Binaphthol Derivatives by Oxidative Coupling of Naphthol Derivatives Catalyzed by Chiral Diamine-Copper Complexes" J. ORG. CHEM., vol. 64, 1999, pages 2264-2271, XP002348777

- D16: DATABASE CROSSFIRE BEILSTEIN Beilstein Institut zur Förderung der Wissenschafter, Frankfurt am Main, DE; 1995, XP002348720 Database accession no. 7305927 (BRN)
- D17: DATABASE CROSSFIRE BEILSTEIN Beilstein Institut zur Förderung der Wissenschafter, Frankfurt am Main, DE; 1988, XP002348721 Database accession no. 179894 (BRN)
- D18: DATABASE CROSSFIRE BEILSTEIN Beilstein Institut zur Förderung der Wissenschafter, Frankfurt am Main, DE; 1994, XP002348722 Database accession no. 6399577 (BRN)
- D19: DATABASE CROSSFIRE BEILSTEIN Beilstein Institut zur Förderung der Wissenschafter, Frankfurt am Main, DE; 2000, XP002348723 Database accession no. 8412429 (BRN)
- D20: DATABASE CROSSFIRE BEILSTEIN Beilstein Institut zur Förderung der Wissenschafter, Frankfurt am Main, DE; 1988, XP002348724 Database accession no. 17092 (BRN)
- D21: EP 0 564 924 A (MILES INC; BAYER CORPORATION) 13 October 1993 (1993-10-13)

Present claims 1 and 6 relate to an extremely large number of possible compounds, in particular in view of the broad definitions of the groups T and R¹. Support within the meaning of Article 5 PCT is to be found, however, for only a very small proportion of the compounds claimed. In the present case, the claims so lack support, and the application so lacks disclosure, that a meaningful search over the whole of the claimed scope is impossible.

Consequently, the search has been carried out for those parts of the claims which appear to be reasonably supported and disclosed, namely those parts wherein both T and R¹ are reasonably defined, i.e. the compounds wherein T is defined as found in claim 7 and R¹ is defined as found in claim 9.

An attempt to search claims 1 and 6 in their full scope resulted in a very huge number of results (many thousand relevant compounds) so that a meaningful search over the complete scope of claims 1 and 6 cannot reasonably be carried out. Thus the search report can only be considered complete for the compounds and use thereof wherein

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY (SEPARATE SHEET)

International application No.

PCT/IB2005/001140

T and R¹ are defined as indicated above, and the compounds and methods cited against the novelty of claims 1-9 are exemplary only.

2. The subject matter defined in claims 1-10 is not novel (Art. 33(2) PCT). The prior art discloses numerous compounds which fall withun the scope of claims 1-10, cf. the relevant passages cited in the search report.

D1 discloses compounds corresponding to claims 1-9 for the treatment of dementia. The subject matter of claims 13-15 lacks novelty in view of this disclosure. Documents D2 and D3 also disclose the therapeutic use of compounds corresponding to claims 1-9 and thus are novelty destroying for the subject matter of claim 13.

D21 discloses, in general form, compounds corresponding to claims 1-9 and the use thereof for the treatmet of diabetes and inflammatory diseases. The subject matter of claims 1-9 and 12-15 therefore is obvious in view of this disclosure (Art. 33(3) PCT).

The use of compounds as defined in claim 10 as medicaments is not derivable from the cited prior art and may be considered inventive.

3. For the assessment of the present claims 14 and 15 on the question whether they are industrially applicable, no unified criteria exist in the PCT Contracting States. The patentability can also be dependent upon the formulation of the claims. The EPO, for example, does not recognize as industrially applicable the subject-matter of claims to the use of a compound in medical treatment, but may allow, however, claims to a known compound for first use in medical treatment and the use of such a compound for the manufacture of a medicament for a new medical treatment.